

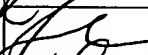

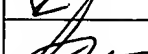

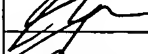


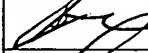





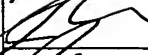




Form PTO/SB/08 INFORMATION DISCLOSURE CITATION IN AN APPLICATION (Use several sheets if necessary)	Docket Number (Optional) BIDM-P01-004	Application Number 10/071,890
	Applicant Frangioni	
	Filing Date February 7, 2002	Group Art Unit 1646

U.S. PATENT DOCUMENTS


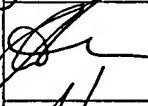

EXAMINER INITIAL	DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE
	AA 5,538,866	7/23/96	Israeli et al.			
	AB 5,804,602	9/8/98	Slusher et al.			
	AC 5,981,209	11/9/99	Slusher et al.			
	AD 6,025,344	2/15/00	Jackson et al.			
	AE 6,071,965	6/6/00	Jackson et al.			
	AF 6,271,245 B1	8/7/01	Jackson et al.			
	AG 6,288,046 B1	9/11/01	Jackson et al.			
	AH 6,348,464 B1	2/19/02	Jackson et al.			
	AI 6,372,726 B1	4/16/02	Slusher et al.			
	AJ 6,384,022 B1	5/7/02	Jackson et al.			
	AK 6,444,657 B1	9/3/02	Slusher et al.			
	AL 6,452,044 B2	9/17/02	Jackson et al.			



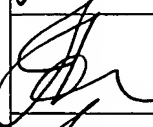


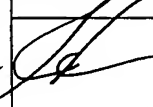
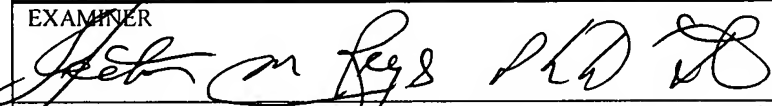
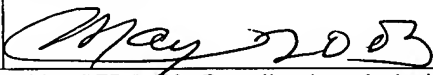
FOREIGN PATENT DOCUMENTS

	DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUBCLASS	Translation	
						YES	NO
	AM WO 97/48399	12/24/97	PCT				
	AN WO 97/48400	12/24/97	PCT				
	AO WO 98/13046	4/2/98	PCT				
	AP WO 98/45257	10/15/98	PCT				
	AQ WO 99/32887	7/1/99	PCT				
	AR WO 02/057222	7/25/02	PCT				

OTHER DOCUMENTS

(Including Author, Title, Date, Pertinent Pages Etc.)

	AS	Berger, U.V. & Schwab, M.E. N-Acetylated Alpha-Linked Acidic Dipeptides may be Involved in Axon-Schwann Cell Signaling. <i>J. Neurocytology</i> 25, 499-512 (1996).
	AT	Jackson, P.F. et al. Design, Synthesis, and Biological Activity of a Potent Inhibitor of the Neuropeptidase N-Acetylated Alpha-Linked Acidic Dipeptidase. <i>J. Med. Chem.</i> 39, 619-622 (1996).
	AU	Kozikowski, A.P. et al. Design of Remarkably Simple, Yet Potent Urea-Based Inhibitors of Glutamate Carboxypeptidase II (NAALADase). <i>J. Med. Chem.</i> 44, 298-301 (2001).

Form PTO/SB/08 INFORMATION DISCLOSURE CITATION IN AN APPLICATION (Use several sheets if necessary)		Docket Number (Optional) BIDM-P01-004	Application Number 10/071,890
		Applicant Frangioni	
		Filing Date February 7, 2002	Group Art Unit 1646
	AV	Lange, P.H. ProstaScint Scan for Staging Prostate Cancer. <i>Urology</i> 57, 402-406 (2001).	
	AW	Nan, F. et al. Dual Function Glutamate-Related Ligands: Discovery of a Novel, Potent Inhibitor of Glutamate Carboxypeptidase II Possessing mGluR3 Agonist Activity. <i>J. Med. Chem.</i> 43, 772-774 (2000).	
	AX	Neale, J.H. et al. N-Acetylaspartylglutamate: The Most Abundant Peptide Neurotransmitter in the Mammalian Central Nervous System. <i>J. Neurochem.</i> 75, 443-452 (2000).	
	AY	Slusher, B.S. et al. Immunocytochemical Localization of the N-Acetyl-Aspartyl-Glutamate (NAAG) Hydrolyzing Enzyme N-Acetylated Alpha-Linked Acidic Dipeptidase (NAALADase). <i>J. Compar. Neuro.</i> 315, 217-229 (1992).	
	AZ	Stauch, B.L. et al. The effects of N-acetylated alpha-linked acidic dipeptidase (NAALADase) inhibitors on [³ H]NAAG catabolism in vivo. <i>Neurosci. Letters</i> 100, 295-300 (1989).	
	BA	Subasinghe, N. et al. Synthesis of Acyclic and Dehydroaspartic Acid Analogues of Ac-Asp-Glu-OH and Their Inhibition of Rat Brain N-Acetylated Alpha-Linked Acidic Dipeptides (NAALADipeptidase). <i>J. Med. Chem.</i> 33, 2734-2744 (1990).	
EXAMINER		DATE CONSIDERED	
			
EXAMINER: Initial if citation considered, whether or not citation is in conformance with MPEP § 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to the applicant.			

Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE




Electronic Information Disclosure Statement

Modified PSMA ligands and uses related thereto

RECEIVED

FEB 19 2003

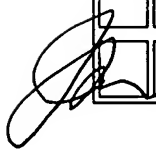

TECH CENTER 1600/2900

Application: 
10/071890
Confirmation: 8688
Applicant(s): John Frangioni
Docket Number: BIDM-P01-004
Group Art Unit: 1646
Examiner:
search string: (6025345).pn.

■ *That each item of information contained in the information disclosure statement was cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement.*

US Patent Documents

Note: Applicant is not required to submit a paper copy of cited US Patent Documents

init	Citation No.	Patent Number	Date	Bar Code	Patentee	Class	Subclass
	P01	6025345	2000-02-15		Jackson et al.		

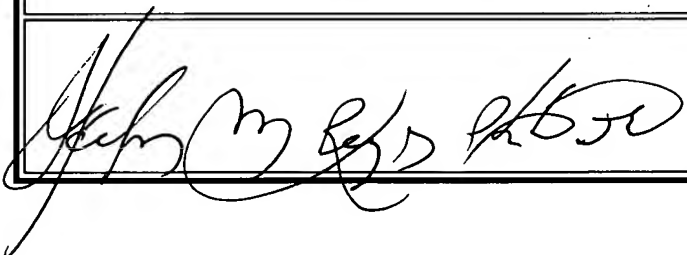
Remarks

(Remarks are not for responding to an office action.)

While the information and references disclosed in this Information Disclosure Statement may be

"material" pursuant to 37 CFR 1.56, it is not intended to constitute an admission that any patent, publication or other information referred to therein is "prior art" for this invention unless specifically designated as such. In accordance with 37 CFR 1.97(g), the filing of this Information Disclosure Statement shall not be construed to mean that a search has been made or that no other material information as defined in 37 CFR 1.56(a) exists. Applicants further reserve the right to take appropriate action to establish the patentability of the disclosed invention over the listed documents should one or more of the documents be applied against the claims of the current application. The Commissioner is hereby authorized to charge any deficiency in the fees filed, asserted to be filed or which should have been filed herewith (or with any paper hereafter filed in this application by this firm) to our Deposit Account No. 18-1945, under Order No. BIDM-P01-004.

Signature

Examiner Name	Date
	June 2003